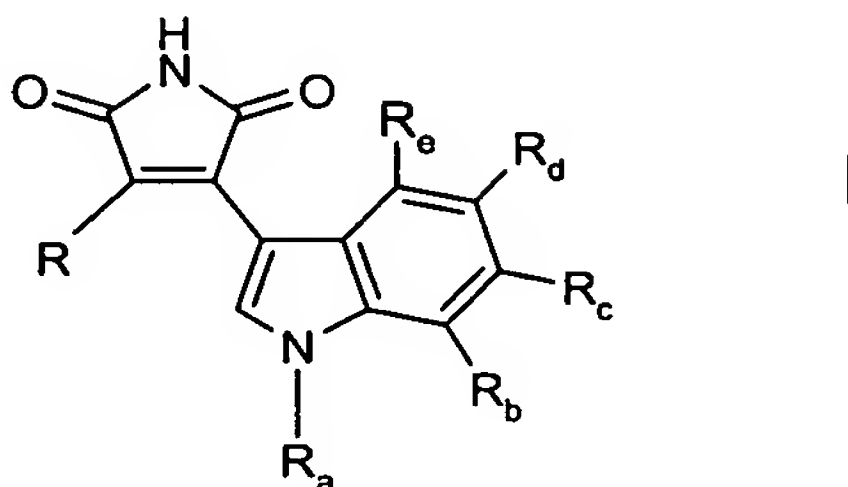


### Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

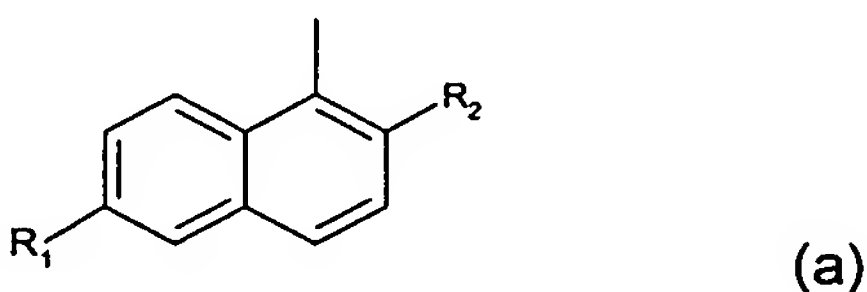
### Listing of Claims:

Claim 1. (Original): A compound of formula I



wherein

R<sub>a</sub> is H; C<sub>1-4</sub>alkyl; or C<sub>1-4</sub>alkyl substituted by OH, NH<sub>2</sub>, NHC<sub>1-4</sub>alkyl or N(di-C<sub>1-4</sub>alkyl)<sub>2</sub>;  
one of R<sub>b</sub>, R<sub>c</sub>, R<sub>d</sub> and R<sub>e</sub> is halogen; C<sub>1-4</sub>alkoxy; or C<sub>1-4</sub>alkyl; and the other three substituents are H; or R<sub>b</sub>, R<sub>d</sub> and R<sub>e</sub> are all H; and  
R is a radical of formula (a)



wherein

R<sub>1</sub> is -(CH<sub>2</sub>)<sub>n</sub>-NR<sub>3</sub>R<sub>4</sub>, wherein  
each of R<sub>3</sub> and R<sub>4</sub>, independently, is H or C<sub>1-4</sub>alkyl; or R<sub>3</sub> and R<sub>4</sub> form together with the nitrogen atom to which they are bound a heterocyclic residue;  
n is 0, 1 or 2; and  
R<sub>2</sub> is H; halogen; C<sub>1-4</sub>alkyl; CF<sub>3</sub>; OH; SH; NH<sub>2</sub>; NO<sub>2</sub>; C<sub>1-4</sub>alkoxy; C<sub>1-4</sub>alkylthio; NHC<sub>1-4</sub>alkyl; N(di-C<sub>1-4</sub>alkyl)<sub>2</sub> or CN;  
or a salt thereof.

Claim 2. (Original): A compound according to claim 1 wherein R<sub>a</sub> is H or methyl; one of R<sub>b</sub>, R<sub>c</sub>, R<sub>d</sub> and R<sub>e</sub> is methyl or ethyl and the other three substituents are H; or R<sub>b</sub>, R<sub>c</sub>, R<sub>d</sub> and R<sub>e</sub> are all H; R<sub>2</sub> is H; Cl, methyl or NO<sub>2</sub>; n is 1; and each of R<sub>3</sub> and R<sub>4</sub>, independently, is H, methyl, ethyl or *i*-propyl; or R<sub>3</sub> and R<sub>4</sub> form together with the nitrogen atom to which they are bound a heterocyclic residue, or a salt thereof.

Claim 3. (Currently amended): A compound according to claim 1 ~~or 2~~ which is selected from 3-(2-Chloro-6-dimethylaminomethyl-naphthalen-1-yl)-4-(1-methyl-1H-indol-3-yl)-pyrrole-2,5-dione;

3-(2-Chloro 6-methylaminomethyl-naphthalen-1-yl)-4-(1H-indol-3-yl)-pyrrole-2,5-dione;

3-(6-Aminomethyl-naphthalen-1-yl)-4-(1-methyl-1H-indol-3-yl)-pyrrole-2,5-dione;

3-(2-Chloro-6-dimethylaminomethyl-naphthalen-1-yl)-4-(1H-indol-3-yl)-pyrrole-2,5-dione;

3-(2-Chloro-6-dimethylaminomethyl-naphthalen-1-yl)-4-(7-methyl-1H-indol-3-yl)-pyrrole-2,5-dione;

3-(2-Chloro-6-methylaminomethyl -naphthalen-1-yl)-4-(7-methyl-1H-indol-3-yl)-pyrrole-2,5-dione;

3-(6-Aminomethyl-naphthalen-1-yl)-4-(1H-indol-3-yl)-pyrrole-2,5-dione;

3-(6-Aminomethyl-naphthalen-1-yl)-4-(7-methyl-1H-indol-3-yl)-pyrrole-2,5-dione; or a salt thereof.

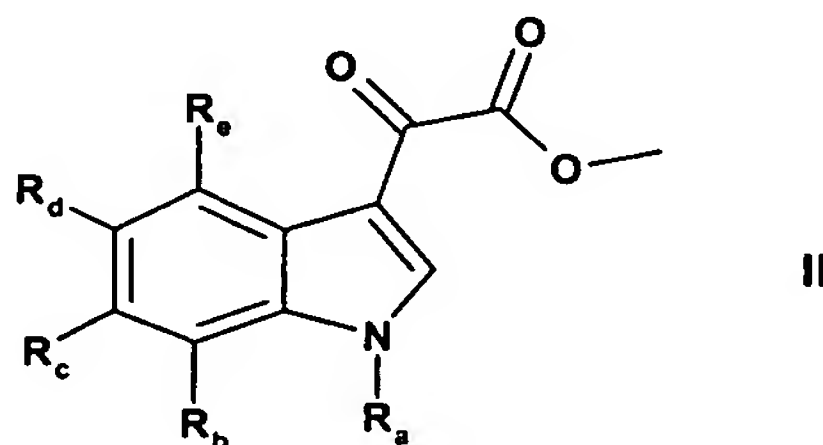
Claim 4. (Currently amended): A compound according to ~~any one of claim 1 to 3~~, in free form or in a pharmaceutically acceptable salt form, for use as a pharmaceutical.

Claims 5. and 6. (Canceled)

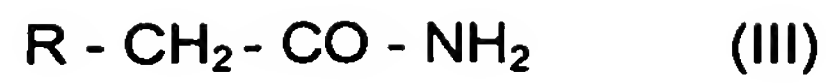
Claim 7. (Currently amended): Use of a compound according to ~~any one of claim 1 to 3~~, in free form or in a pharmaceutically acceptable salt form, or a pharmaceutical composition according to claim 5 in the manufacture of a medicament for treatment and/or prevention of T-cell mediated acute or chronic inflammatory diseases or disorders, autoimmune diseases, graft rejection, cancer or infectious diseases.

Claim 8. (Currently amended): A pharmaceutical combination comprising a compound according to ~~any one of claim 1 to 3~~, in free form or in a pharmaceutically acceptable salt form, and a further agent selected from immunosuppressant, immunomodulatory, anti-inflammatory, chemotherapeutic, antiproliferative and anti-diabetic agents.

Claim 9. (Currently amended): A process for the production of the compound of formula I according to claim 1 ~~or claim 2~~, which process comprises reacting a compound of formula II



wherein R<sub>a</sub> ; R<sub>b</sub>; R<sub>c</sub>, R<sub>d</sub> and R<sub>e</sub> are as defined in claim 1 ~~and claim 2~~,  
with a compound of formula III



wherein R is as defined in claim 1 and claim 2,  
and, where required, converting the resulting compound of formula I obtained in free form to a salt form or vice versa, as appropriate.

Claim 10. (Currently amended): A method for treating or preventing disorders or diseases mediated by T lymphocytes and/or PKC, in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound according to any one of claim 1 to 3, or a pharmaceutically acceptable salt thereof.